

RECEIVED SEHQ-0902-15201

Product Safety and Regulatory Affairs

2002 SEP 24 AM 6: 18

62219

September 13, 2002

Document Processing Center (TS-790) Office of Toxic Substances US Environmental Protection Agency 401 M Street SW Washington, DC 20460

Attn: TSCA 8(e) Coordinator

Dear Sir or Madam:

Please find enclosed confidential and non-confidential copies of a Section 8(e) submission relevant to testing we have performed on one of our products. Please hold the trade name and indicated compositional information confidential.

Sincerely,

Alan Taylor, Ph.D.

Director Product Safety and Regulatory Affairs

Crompton Como ation

8 EHQ-02-15201 45020000127





RECEIVED OPPT CBIC

2002 SEP 24 AM 6: 18

Product Safety & Regulatory Affairs

3500 South State Route 2 Friendly, WV 26146 Phone: (304) 652-8000

Fax: (304) 652-1478

Document Processing Center (TS-790) Office of Toxic Substances US Environmental Protection Agency 401 M Street SW Washington, DC 20460

Attn: TSCA 8(e) Coordinator

September 13, 2002

Dear Sir or Madam:

Crompton Corporation herewith submits the following information pursuant to TSCA Section 8(e) concerning preliminary audited information from an acute oral toxicity (gavage) study with rats on a silyl ketene acetal known commercially as ******.

***** is a mixture containing a maximum concentration of the following:

Concentration (wt. %)	Chemical Name	CAS#
	3,5-Dioxa-2,6-disilaheptane, 2,2,6,6-tetramethyl-4-(1-methylethylidene)-	31469-25-7

The information provided in this letter is based on the instructions regarding reportability of toxicity data stated in the EPA's <u>TSCA Section 8(e) Reporting Guide</u>. This information was received as described herein on September 3, 2002. This information is summarized below.

A study of ***** in Sprague-Dawley rats was conducted in order to evaluate the possible acute toxic effects of ***** when administered once orally (gavage) to rats. The oral route was selected since this is an acceptable and standard method for administering the test article under OECD Guideline 407. Ingestion is not a potential route of exposure for humans, other than through accidental ingestion by workers handling the test article.

Treatment was by oral gavage with the neat test substance. The calculated LD_{50} for ****** was greater than 2000 mg/kg of body weight. There were no mortalities. Clinical signs were observed and included impaired muscle coordination in all animals on day 0 and persisted in all males on day 1, but had subsided by day 2. Prostration was observed in three males on day 0 and in one male on day 2. Hypoactivity was observed in four males and one female on day 0 and in one male on day 1. Other signs included various discolored areas due to discharges/excretions, increased respiration rate (one animal), labored respiration and partial closure of the eye(s) (one animal). All animals appeared normal by day 4.

Please contact the undersigned with questions, if any, at (203) 573-2219.

Sincerely,

Alan Taylor, Ph.D.

Director, Product Safety and Regulatory Affairs

Crompton Corporation

Sher Says